

APR 06 2006

Sheet 1 of 5

FORM PTO-101 (Rev. 2-32) U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE  INFORMATION DISCLOSURE STATEMENT BY APPLICANT  (Use several sheets if necessary)	ATTY. DOCKET NO. 30307A-DIV1	SERIAL NO. 10/876,297
	APPLICANT: KNOLL, Joan et al	
	FILING DATE: 6/23/04	GROUP:

## U.S. PATENT DOCUMENTS

EXAM INITIAL	U.S.	DOCUMENT NUMBER							INVENTOR NAME	CLASS	SUB-CLASS	ISSUE DATE (PATENT); PUBLICATION DATE (PUBLISHED APPLICATION); OR FILING DATE (NON-PUBLISHED APPLICATION)
SP	2002/	0	1	9	2	6	9	2	Palanisamy et al	435/6		12/2002
SP		6	1	2	1	4	1	9	Rowley et al	530/350		09/2000
SP		5	4	4	7	8	4	1	Gray et al			9/1995
SP		5	7	5	6	6	9	6	Gray et al			5/1998
SP		5	7	2	1	0	9	8	Pinkel et al			2/1998
SP		6	2	2	2	0	2	9	Edwards et al			4/2001
SP		5	8	1	1	2	3	1	Farr et al			9/1998
SP		6	0	4	0	1	4	0	Croce et al			3/2000

## FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER							PUBLICATION DATE	COUNTRY	CLASS	SUB-CLASS	TRANSLATION	
													YES	NO

## OTHER DOCUMENTS (Including Publisher, Author, Title, Relevant Pages, and Date and Place of Publication)

SP			Tanaka et al., Cancer Genetics and Cytogenetics, 1999. 133: 29-35.
SP			Rockman et al., Australian Journal of Medical Science, May 15, 1994; 56-57.
SP			DeRisi et al., Nature Genetics, December 14, 1996; 457-460
SP			Sakuma, GenBank Accession No. AB001872, Feb. 13, 1998
SP			Urano, GenBank Accession No. AB002058, August 29, 1997

/Steven Pohnert/ (08/30/2006)

EXAMINER: Initial if citation considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

FORM PTO-1449 U.S. DEPARTMENT OF COMMERCE (Rev. 2-32) PATENT AND TRADEMARK OFFICE  INFORMATION DISCLOSURE STATEMENT BY APPLICANT  (Use several sheets if necessary)	ATTY. DOCKET NO. 30307A-DIV1	SERIAL NO. 10/876,297	
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	FILING DATE: 6/23/04	GROUP:	CONF. NO:

**U.S. PATENT DOCUMENTS**

[illegible]

## FOREIGN PATENT DOCUMENTS

[illegible]

**OTHER DOCUMENTS (Including Publisher, Author, Title, Relevant Pages, and Date and Place of Publication)**

SP		Inoue et al., GenBank Accession No. AB002135, July 23, 1999
SP		Watanabe, GenBank Accession No. AB003723, Feb. 25, 1998
SP		Kamei, GenBank Accession No. AB003592, July 19, 1999
SP		Ohno, GenBank Accession No. AB000114, Feb. 5, 1999
SP		Shimomura, GenBank Accession No. AB0000095, March 4, 1998

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	FILING DATE: 6/23/04	GROUP:	CONF. NO:

## U.S. PATENT DOCUMENTS

[illegible]

## FOREIGN PATENT DOCUMENTS

[illegible]

**OTHER DOCUMENTS (Including Publisher, Author, Title, Relevant Pages, and Date and Place of Publication)**

SP		Satoh, GenBank Accession No. AB000276, November 6, 1997
SP		Yokouchi, GenBank Accession No. AB000520, September 26, 1997
SP		Fununaga, GenBank Accession No. AB000409, Feb. 5, 1999
SP		Fukuta, GenBank Accession No. AB003791, Feb. 14, 1998
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SP		Ishihara, GenBank Accession No. AB003333, Feb. 26, 1999
SP		11:000-000 2001 by Cold Spring Harbor Laboratory Press ISSN 1088-9051/01; Sequence-Based Design of Single-Copy Genomic DNZ Probes for Fluorescence In Situ Hybridization; pgs. 1-9
SP		Med-Genet 1999; 36: 657-663; Monosomy 1P36; Slavotinek, Shaffer, Shapira
SP		Current Opinion in Structural Biology, 1998, 8: Repeats in genomic DNA; mining and meaning; Jerzy Jurka; 333-337
SP		Genomics 29; 397-402 (1995); Human Genomic characterization of a Novel Locus-Specific Repetitive Sequence

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**U.S. PATENT DOCUMENTS**

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## FOREIGN PATENT DOCUMENTS

[illegible]

**OTHER DOCUMENTS (Including Publisher, Author, Title, Relevant Pages, and Date and Place of Publication)**

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SP		Molecular Cloning, A Laboratory Manual, Second Edition, 1989 by Cold Spring Harbor Laboratory Press; The Effects of Length and Degeneracy of the Oligonucleotide on the Specificity of Hybridization.
SP		Science 1990 Oct 5; 250(4977): 94-8; Chromosomal region of the cystic fibrosis gene in yeast artificial chromosomes; a model for human genome mapping; Green Ed, Olson MV.

<b>FORM PTO-1449</b> (Rev. 2-32)	<b>U.S. DEPARTMENT OF COMMERCE</b> <b>PATENT AND TRADEMARK OFFICE</b>	<b>ATTY. DOCKET NO.</b> 33026	<b>SERIAL NO.</b>
		<b>APPLICANT:</b> ROGAN, PETER et al.	
		<b>FILING DATE:</b> September 30, 2003	<b>GROUP:</b>

## U.S. PATENT DOCUMENTS

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## FOREIGN PATENT DOCUMENTS

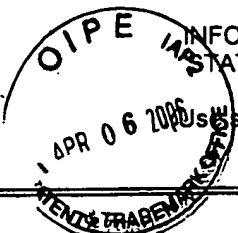
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**OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)**

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SP			Carter et al., Comparative Analysis of Comparative Genomic Hybridization Microarray Technologies: Report of a Workshop Sponsored by the Wellcome Trust

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FORM PTO-1449 (Rev. 2-32)	U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTY. DOCKET NO. 33026	SERIAL NO. 10/676,248
 INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use several sheets if necessary)		APPLICANT: ROGAN et al	
		FILING DATE: September 30, 2003	GROUP: 1645

## U.S. PATENT DOCUMENTS

EXAM. INITIAL		DOCUMENT NUMBER							DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
SP		6	0	0	7	9	9	4	12/28/1999	WARD et al			
SP		6	4	0	0	0	3	3	8/8/2000	SMITH et al			
SP		6	4	0	6	8	2	0	6/18/2002	VOLKERS et al			
SP		6	5	2	1	4	2	7	02/18/03	EVANS			
SP	0	9	5	7	3	0	8	0	05/16/00	KNOLL			
SP	0	9	8	5	4	8	6	7	05/14/01	KNOLL			

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SP	WO	0	1	8	8	0	8	9	11/22/2001				YES	NO
SP	WO	9	9	0	6	5	5	2	2/11/99	PCT				

## OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

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SP		FISHing for mechanisms of cytogenetically defined terminal deletions using chromosome-specific subtelomeric probes; Ballif, Blake C., Kashork, Catherine D., and Schaffer, Lisa G.; European Journal of Human Genetics (2000) B, pp 764-770
SP		Identification of Cryptic Rearrangements in Patients with 18q- Deletion Syndrome; Brkanac, Zoran; Cody, Jannine D., Leach, Robin J., and DuPont, Barbara R.; American Journal of Human Genetics 62: 1998; pp 1500-1506
SP		Structure and Polymorphism of Human Telomere-Associated DNA; Brown, MacKinnon, Villasante, Spurr, Buckle and Dobson; Cell; pp 119-132
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SP		Maternal Balanced Translocation Leading to Partial Duplication of 4q and Partial Deletion of 1p Band-Specific Painting Probes Generated by Chromosome Microdissection; Chen, Grebe, Guan, Notohamiprodjo, Nutting, Stone, Trent and Sandberg; American Journal of Medical Genetics 71 (1997) pp 160-166
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SP		Clinical phenotype associated with terminal 2q37 deletion; Conrad, Dewald, Christensen, Lopez, Higgins, and Pierpont; Clinical Genetics 1995; Vol 48; pp 134-139
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SP		Cryptic 6q Subtelomeric Deletion Associated With a Paracentric Inversion in a Mildly Retarded Child; Lorda-Sanchez, Pajares, Roche, Sanz, de Alba, Gonzalez-Gonzalez, Ibanez, Ramos, and Ayuso; American Journal of Medical Genetics; Vol. 95 (2000) pp 336-338
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SP		Chromosome 1p36 Deletions: The Clinical Phenotype and Molecular Characterization of a Common Newly Delineated Syndrome; Shapira, McCaskill, Northrup, Spikes, Elder, Sutton, Korenberg, Greenberg, and Shaffer; American Journal of Human Genetics; Vol 61 (1997) pp 642-650



